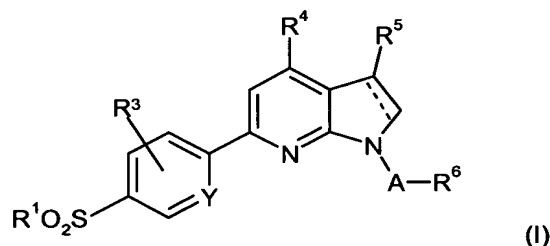


In the Claims:

Please amend the claims as follows:

1. (Original) A compound of formula (I)



or a pharmaceutically acceptable salt thereof in which:

Y is selected from the group consisting of CH or nitrogen;

R¹ is selected from the group consisting of C₁₋₆alkyl, NH₂ and R²CONH;

R² is selected from the group consisting of H, C₁₋₆alkyl, C₁₋₆alkoxy, C₁₋₆alkyloxyC₁₋₆alkyl, phenyl, HO₂CC₁₋₆alkyl, C₁₋₆alkyloxyCOOC₁₋₆alkyl, C₁₋₆alkyloxyCO, H₂NC₁₋₆alkyl, C₁₋₆alkyloxyCONHC₁₋₆alkyl and C₁₋₆alkylCONHC₁₋₆alkyl;

R³ is selected from the group consisting of H and halogen;

R⁴ is selected from the group consisting of H, C₁₋₅alkyl, and C₁₋₂alkyl substituted by one to five fluorine atoms;

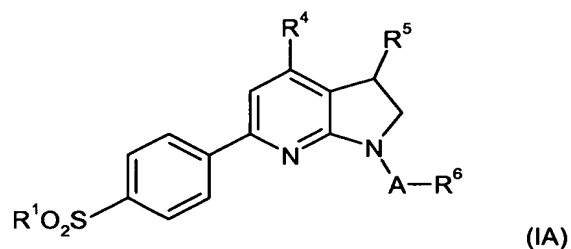
R⁵ is selected from the group consisting of H, CHO, and C₁₋₆alkyl which is unsubstituted or is substituted one or more times by halogen or hydroxy;

A is $(\text{CH}_2)_n$ or $-\text{SO}_2-$;

R⁶ is selected from the group consisting of C₁₋₆alkyl, C₄₋₈ cycloalkyl, phenyl and 6-membered heteroaryl, wherein the phenyl and 6-membered heteroaryl ring may be unsubstituted or substituted one or more times by halogen or C₁₋₆ alkyl; and
n is 0 to 3.

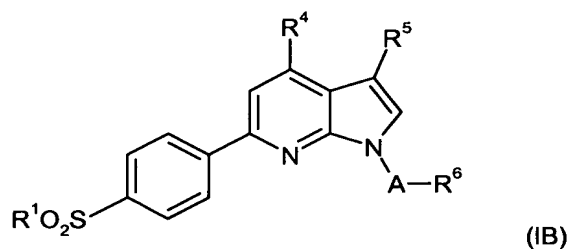
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2. (Original) A compound of formula (IA)



or a pharmaceutically acceptable salt thereof, in which all substituents are as for a compound of formula (I) as defined in claim 1.

3. (Original) A compound of formula (IB)



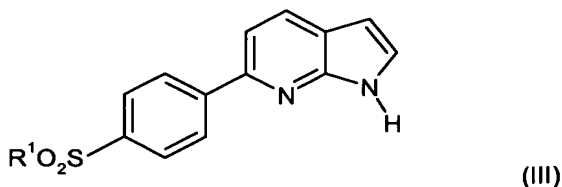
or a pharmaceutically acceptable salt thereof, in which all substituents are as for a compound of formula (I) as defined in claim 1.

4. (Currently Amended) A compound according to ~~any of~~ claims 1 ~~to~~ 3 wherein R^1 is C_{1-6} alkyl.
5. (Currently Amended) A compound according to ~~any of~~ claims 1 ~~to~~ 4 wherein R^4 is H, CHF_2 , CH_2F , CF_3 or C_{1-4} alkyl.
6. (Currently Amended) A compound according to ~~any of~~ claims 1 ~~to~~ 5 wherein R^5 is H, C_{1-4} alkyl, $-CHO$, or $-(CH_2)_nCH_2OH$.

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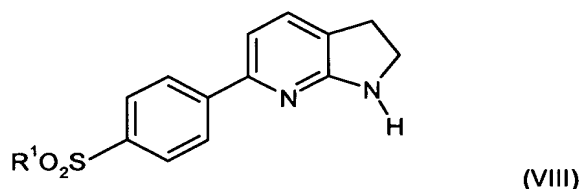
7. (Currently Amended) A compound according to ~~any of~~ claims 1 ~~to~~ 6 wherein R⁶ is C₃₋₅alkyl, cyclohexyl, pyridyl optionally substituted by C₁₋₃alkyl, or phenyl optionally substituted by halogen.
8. (Currently Amended) A compound according to ~~any of~~ claims 1 ~~to~~ 7 wherein n is 0 or 1.
9. (Original) A compound according to claim 3 wherein R¹ is C₁₋₃alkyl, R⁴ is H, CHF₂, CH₂F, CF₃ or C₁₋₄alkyl, R⁵ is H, C₁₋₄alkyl, -CHO, or -CH₂OH, n is 1, and R⁶ is C₃₋₅alkyl, cyclohexyl, pyridyl optionally substituted by C₁₋₃alkyl, or phenyl optionally substituted by halogen.
10. (Original) A compound according to claim 3 wherein R¹ is C₁₋₃alkyl, R⁴ is H, CHF₂, CH₂F, CF₃ or C₁₋₄alkyl, R⁵ is H, C₁₋₄alkyl, -CHO, or -CH₂OH, n is 0, and R⁶ is phenyl optionally substituted by halogen.
11. (Original) A compound according to claim 3 wherein R¹ is CH₃, R³ is H, R⁴ is H, R⁵ is H, C₁₋₄alkyl, -CHO, or -CH₂OH, A is (CH₂)_n and n is 1, and R⁶ is C₃₋₅alkyl, cyclohexyl, pyridyl optionally substituted by CH₃, or phenyl optionally substituted by chloro.
12. (Original) A compound according to claim 3 wherein R¹ is CH₃, R³ is H, R⁴ is H, R⁵ is H, A is (CH₂)_n and n is 0, and R⁶ is phenyl optionally substituted by fluoro.
13. Canceled.
14. (Original) A process for the preparation of compounds of formula (IA), as defined in claim 2, where each of R⁴ and R⁵ is hydrogen, which comprises:

reducing a compound of formula (III)

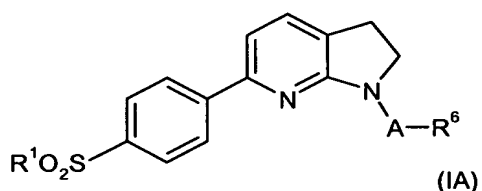


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to form a compound of formula (VIII);



reacting said compound of formula (VIII) with a compound R^6 -A-X, or a protected derivative thereof, where X is a halogen, such as Cl, Br or I, or a sulfonate such as methanesulfonate, (4-methyl)benzenesulfonate or trifluoromethanesulfonate, and A and R^6 are as hereinbefore defined; such as to produce a compound of formula (IA), wherein R^4 and R^5 are both hydrogen

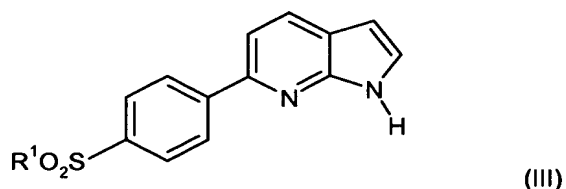


and thereafter and if necessary,
interconverting said compound of formula (IA) into another
compound of formula (IA); and/or

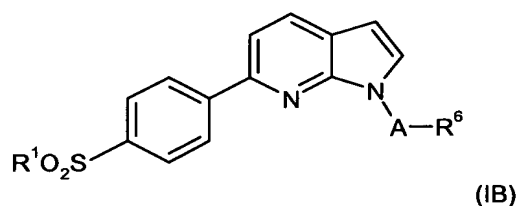
deprotecting a protected derivative of compound of formula (IA).

15. (Original) A process for the preparation of compounds of formula (IB), as defined in claim 3, where each of R^4 and R^5 is hydrogen, which comprises:

reacting a compound R^6 -A-X (II) or a protected derivative thereof,
with a compound of formula (III)



where X is a halogen, such as Cl, Br or I, or a sulfonate, such as methanesulfonate, (4-methyl)benzenesulfonate or trifluoromethanesulfonate, and R⁶ and A are as hereinbefore defined, to produce a compound of formula (IB) in accordance with the present invention :



and thereafter and if necessary,
interconverting said compound of formula (IB) into another
compound of formula (I); and/or

deprotecting a protected derivative of compound of formula (IB).

16. (Currently Amended) A pharmaceutical composition comprising a compound of formula (I) as defined in ~~any of claims 1 to 10~~ in admixture with one or more physiologically acceptable carriers or excipients.
17. (Currently Amended) A compound of formula (I) as defined in ~~any of claims 1 to 10~~ for use in human or veterinary medicine.
18. (Currently Amended) A method of treating a human or animal subject suffering from a condition which is mediated by COX-2 which comprises administering to said subject an effective amount of a compound of formula (I) as defined in ~~any of claims 1 to 10~~.

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19. (Currently Amended) A method of treating a human or animal subject suffering from an inflammatory disorder, which method comprises administering to said subject an effective amount of a compound of formula (I) as defined in ~~any of claims 1 to 10~~.
20. (Currently Amended) The use of a compound of formula (I) as defined in ~~any of claims 1 to 10~~ for the manufacture of a therapeutic agent for the treatment of a condition which is mediated by COX-2.
21. (Currently Amended) The use of a compound of formula (I) as defined in ~~any of claims 1 to 10~~ for the manufacture of a therapeutic agent for the treatment of an inflammatory disorder.